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Drug-induced Apoptosis in Hepatoma Cells Is Mediated by the CD95 (APO-1/Fas) Receptor/Ligand System and Involves Activation of Wild-type p53

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Abstract

Chemotherapeutic drugs are cytotoxic by induction of apoptosis in drug-sensitive cells. We investigated the mechanism of bleomycin-induced cytotoxicity in hepatoma cells. At concentrations present in the sera of patients during therapy, bleomycin induced transient accumulation of nuclear wild-type (wt) p53 and upregulated expression of cell surface CD95 (APO-1/Fas) receptor in hepatoma cells carrying wt p53 (HepG2). Bleomycin did not increase CD95 in hepatoma cells with mutated p53 (Huh7) or in hepatoma cells which were p53^{-/-} (Hep3B). In addition, sensitivity towards CD95-mediated apoptosis was also increased in wt p53 positive HepG2 cells. Microinjection of wt p53 cDNA into HepG2 cells had the same effect. In contrast, bleomycin did not enhance susceptibility towards CD95-mediated apoptosis in Huh7 and in Hep3B cells. Furthermore, bleomycin treatment of HepG2 cells increased CD95 ligand (CD95L) mRNA expression. Most notably, bleomycin-induced apoptosis in HepG2 cells was almost completely inhibited by antibodies which interfere with CD95 receptor/ligand interaction. These data suggest that apoptosis induced by bleomycin is mediated, at least in part, by p53-dependent stimulation of the CD95 receptor/ligand system. The same applies to other anti-cancer drugs such as cisplatin and methotrexate. These data may have major consequences for drug treatment of cancer and the explanation of drug sensitivity and resistance. (*J. Clin. Invest.* 1997; 99:403–413.) Key words: apoptosis • CD95 • p53 • cancer therapy • drug sensitivity

Introduction

Chemotherapeutic drugs cause DNA damage and kill cancer cells mainly by apoptosis (1, 2). The mechanisms of drug-induced apoptosis, however, are poorly understood. Recent studies suggest that p53 may be required for apoptosis in cells treated with chemotherapeutic drugs or radiation (1, 3). The p53 tumor suppressor gene is an essential mediator of the cel-

lular response to DNA damage in mammalian cells. Both G₁ cell cycle arrest (4–6) and apoptosis (7–9) after DNA damage were shown to depend on normal p53 function (10). The p53 protein senses genotoxic stress or DNA damage. This results in nuclear accumulation of p53. Subsequently, p53 activates the transcription of several genes whose products are involved in DNA repair or apoptosis. Some of the genes regulated by p53 have been identified. These genes include muscle creatinine kinase (11), gadd45, waf-1/cip-1, bcl2/bax, cyclin G, mdm2, TGF-α, IGF-binding protein, and Fas. (4–6, 12–15). The exact mediators of apoptosis after p53 activation are not known. Owen-Schaub et al. (16) have shown that introduction of wt or temperature-sensitive p53 into human tumor cell lines enhances expression of the CD95 (APO-1/Fas) receptor.

CD95 is a type I transmembrane receptor that belongs to the nerve growth factor/tumor necrosis factor receptor superfamily. Triggering of CD95 by the CD95 ligand (CD95L) or by agonistic antibodies results in rapid induction of apoptosis in susceptible cells (17–19). CD95L has recently been cloned and found to be a type II transmembrane protein that belongs to the corresponding TNF family (20, 21).

The CD95 receptor/ligand system plays an important role in downregulation of the immune response. In addition, the CD95 and the perforin system mediate T cell cytotoxicity (22–24). CD95 is constitutively expressed in a wide range of tissues, including the liver (25–27). The liver is highly sensitive to CD95-mediated induction of apoptosis. Thus, mice injected intraperitoneally with anti-CD95 antibody died within several hours from fulminant hepatic failure due to generalized apoptosis of hepatocytes (26). Likewise, human hepatocytes in primary culture undergo apoptosis upon stimulation with agonistic anti-CD95 antibodies (25). Moreover, upregulated CD95 receptor- and/or CD95L mRNA levels were detected in different acute and chronic liver diseases (25).

We have analyzed the mechanisms by which cytotoxic drugs kill hepatoma cells in vitro. We found that these drugs kill these cells by induction of apoptosis. Therefore, we tested whether drug-induced apoptosis involves the CD95 receptor/ligand system and is dependent on the function of p53. Our data show that antineoplastic drugs induce apoptosis in wt p53 carrying hepatoma cells by upregulation of the CD95 receptor and the CD95L.

Methods

Cell lines

The following cell lines were used: (a) HepG2 cells derived from a human hepatoblastoma (28) only expressing small amounts of wild-type p53 (unpublished sequence data from our laboratory); (b) Huh7 cells derived from a hepatocellular carcinoma (29) and shown to ex-

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press p53 with increased half life as a result of a point mutation at codon 220 (30); and (c) Hep3B cells (28) deficient of p53 (31). HepG2 and Huh7 cells were maintained in Dulbecco's modified Eagle medium (DMEM; Flow Laboratories, Meckenheim, Germany) containing 10% FCS, 5 mmol/liter glutamine, 100 µg/ml penicillin (all from Flow Laboratories).

Hep3B were grown in MEM (Milieu Essentiel Minimum; Laboratoires Eurobio, Raunheim, Germany).

Treatment with anticancer agents

The different cell lines were treated with bleomycin (Mack, Illertissen, Germany) at a dose range of 60 ng/ml to 6 mg/ml, or with doxorubicin (Rhône-Poulenc, Köln, Germany) at a dose range of 0.4 ng/ml to 4 µg/ml for 3 h up to 72 h. Furthermore, cisplatin at a dose range of 0.1 ng/ml to 0.1 mg/ml or methotrexate at a dose range of 0.1 ng/ml to 1 mg/ml, were applied. The concentrations relevant for therapy are 1.5–3.0 µg/ml for bleomycin (32), 0.4–1.6 µg/ml for cisplatin (33, 34), 7–12 µg/ml for methotrexate (35), and 0.001–0.02 µg/ml for doxorubicin (36) in patients' sera.

Treatment with IgG 3 anti-APO-1 and F(ab)'₂-anti-APO-1

CD95 receptor stimulation was performed with the monoclonal antibody anti-APO-1 at concentrations of 100 ng/ml as described (17, 37, 38). CD95 receptor blocking experiments were performed by treatment with F(ab)'₂-anti-APO-1 fragments as previously described (38).

Statistical analysis

To examine whether synergy (39, 40) between APO-1 stimulation and concurrent bleomycin treatment is observed, a balanced two-way ANOVA (model with fixed effects) was performed. Statistical analysis was carried out with the SAS software system (SAS Institute Inc., Cary, NC).

Immunohistochemistry

HepG2 cells grown on cover slips were fixed for 2 min in a 1:1 (vol/vol) mixture of acetone and methanol and air dried. Immunohistochemistry of p53 was performed as described (41), using Do1 (Dianova, Hamburg, Germany) as first antibody.

Immunoblot (Western blot)

Cultured cells were homogenized in sample buffer (50 mM Tris-Cl, pH 6.8, 2% SDS, 100 mM dithiothreitol). Protein concentration was measured using the Bio-Rad protein assay (Bio-Rad Laboratories, Munich, Germany). Briefly, an aliquot of each sample equivalent to 10 µg protein was boiled in 20 µl sample buffer, separated on a 12% SDS-polyacrylamide gel (PAGE), and electrophoretically transferred to nitrocellulose filters using the Bio-Rad electrotransfer system (Bio-Rad Laboratories, Munich, Germany). P53 was detected with the mouse monoclonal antibody PAb 1801 (Dianova, Hamburg, Germany) or the polyclonal rabbit antiserum HZp53R as first antibody as described (42).

Metabolic labeling with [³⁵S]methionine, immunoprecipitation, and autoradiography

HepG2 and Huh7 cells were plated in six-well dishes (2 × 10⁵/well) at 60% confluence. Cells were treated with bleomycin or doxorubicin as described above for 3 h. HepG2 and Huh7 cells were then metabolically labeled for 3 h in 2 ml of 10% dialyzed FCS/methionine-free medium with 20 µCi/ml [³⁵S]methionine. Cells were collected with a rubber policeman and centrifuged for 5 min at 13,000 g. The cells were lysed in lysis buffer (50 mM Tris, pH 8.0, 150 mM NaCl, 5 mM EDTA, 1% NP-40, 1 mM PMSF).

Monoclonal antibodies PAb 1801 or PAb 421 (Dianova, Hamburg, Germany) were adsorbed to 20 µl of preswollen protein A-Sepharose (Sigma, Munich, Germany) in phosphate-buffered saline (PBS) for 2 h at room temperature. After washing three times with lysis buffer supernatants of the cells were added and incubated 2 h at room temperature.

The immunoprecipitates were boiled in sample buffer and separated by 10% SDS-PAGE. The gel was then dried and subjected to autoradiography.

[³H]Thymidine incorporation

Cell proliferation was determined in 35-mm culture plates. 1 µCi/ml [³H]thymidine (Amersham Buchler, Braunschweig, Germany) was added to each culture well for 3 h before harvesting the cells. Subsequently, the tissue culture plates were placed on ice and the liver cells washed twice with ice-cold PBS and harvested with 1% Triton X-100/PBS.

Incorporated [³H]thymidine was measured in a liquid scintillation counter (Beckman, Munich, Germany). Growth inhibition by bleomycin or doxorubicin was calculated as the percentage reduction in [³H]thymidine uptake compared with untreated controls. Results are shown as the mean of six independent culture plates ± SD.

Detection of apoptosis

FACS[®] analysis. Apoptosis was assessed by fluorescence-activated cell sorting (FACS[®]) analysis carried out in a FACScan[®] flow cytometer (Becton Dickinson, Heidelberg, Germany) using the LYSIS II software system.

Hepatocytes, floating in the culture medium were collected by centrifugation at 200 g. Adherent hepatocytes were harvested by incubation with 1% trypsin for 1 min. The cells were washed with PBS, fixed in 70% ethanol and stained with propidium iodide (50 µg/ml PBS). DNA fluorescence was measured in a FACScan[®] according to the method of Nicoletti et al. (43).

DNA fragmentation. Intermuclosomal DNA fragmentation was assessed by the method of Sellins and Cohen as described (44) after harvesting adherent and floating cells from a 70 cm² subconfluent culture dish.

DAPI and TUNEL staining. Cells were grown on plastic cover slips. DAPI (4',6-diamidino-2-phenylindole; Sigma, Munich, Germany) staining of cellular DNA, was performed as described (25). The cover slips were viewed under the fluorescence microscope (Zeiss, Axiophot, Göttingen, Germany) at 420 nm.

TUNEL staining (45) was performed as described in the Boehringer kit (Boehringer, Mannheim, Germany).

Microinjection

For microinjection about 2,000 HepG2 cells were seeded onto CELLocate coverslips (Eppendorf, Hamburg, Germany). The following day about 50 cells (per coverslip) in DMEM supplemented with 10% FCS, 1× non-essential amino acids (GIBCO, Eggenstein, Germany) and 10 mM Hepes (pH 7.4) were injected with 100 ng/ml H₂O p53 wild-type DNA or different control DNA constructs with the CMV promoter in 1% tetramethylrhodamine isothiocyanate-dextran (M_r = 4400) using the Microinjector 5242 (Eppendorf, Hamburg, Germany) connected to the automated injection system (Zeiss, Göttingen, Germany). The settings were 70–100 hPa and 0.1 s. After 48 h of incubation the medium was changed and 100 ng/ml agonistic IgG3 anti-APO-1 mAb were added and the cells were further incubated for 24 h. Surviving microinjected cells were counted under the fluorescence microscope.

Immunostaining of p53. After microinjection and further incubation for 48 h HepG2 cells were fixed in methanol and washed with PBS/0.1% Triton X-100. p53 was stained with the mouse monoclonal antibody PAb 1801 (10 ng/ml) followed by FITC-conjugated goat anti-mouse IgG (5 ng/ml).

Detection of CD95 (APO-1/Fas)

CD95 cell surface expression was assessed by FACScan[®]. Anti-APO-1 (IgG3, κ) was used as purified biotinylated antibody. Quantum Red streptavidin (Sigma, Munich, Germany) was used as secondary reagent for indirect immunofluorescence. All purchased reagents were used according to the manufacturer's instructions. Fluorescence analysis was carried out at 4°C in PBS with 0.1% sodium azide. Hepatoma

cells were incubated in 200 μ l staining buffer with biotinylated anti-APO-1. After 15 min incubation, cells were washed twice, incubated for 15 min with Quantum Red streptavidin, washed twice again and assayed. Upon data acquisition a gate was set on intact cells by forward/side scatter analysis and 5,000–10,000 viable cells were analyzed.

Detection of CD95 ligand expression by *in situ* hybridization

HepG2 cells at 60% confluence were incubated on two-well chamber slides (Lab-Tek[®], Nunc Inc., Naperville, IL) with 6 mg/ml bleomycin for 24 h. After washing with PBS the cells were fixed for 20 min in 4% paraformaldehyde, dehydrated in ethanol and air dried.

For *in situ* hybridization the cells were pretreated for 20 min at 37°C with proteinase K (1 μ g/ml in 2 \times SSC; 0.1% SDS). The reaction was stopped by 0.1 M glycine, the cells were fixed with 4% paraformaldehyde and washed with 2 \times SSC. After treatment for 10 min in

0.25% acetic anhydride in 0.1 M triethanolamine, pH 8.0 the cells were dehydrated through an ascending series of ethanol and air dried.

Digoxygenin labeled anti-sense RNA corresponding to the sequence coding for amino acids 100 to 246 of the human CD95-ligand was generated using *in vitro* transcription with T7 polymerase (Boehringer, Mannheim, Germany). The length was reduced to approximately 250 nucleotide fragments by limited alkaline hydrolysis (46). The hybridization mixture contained 100 pg/ μ l digoxygenin-labeled RNA, 50% deionized formamide, 10% dextran sulfate, 2 \times SSC, 1 \times Denhardt's mixture (0.02% wt/vol), 0.1% SDS, 500 μ g/ml yeast tRNA, 100 μ g/ml herring sperm DNA. This mixture was spread over the pretreated, dried cells and covered with parafilm. The slides were incubated overnight at 37°C in a moist chamber containing 50% formamide. Thereafter, the slides were washed with 2 \times SSC (30 min), digested with 5 μ g RNase A (20 min) and again washed with 0.5 \times SSC (20 min) at 37°C.

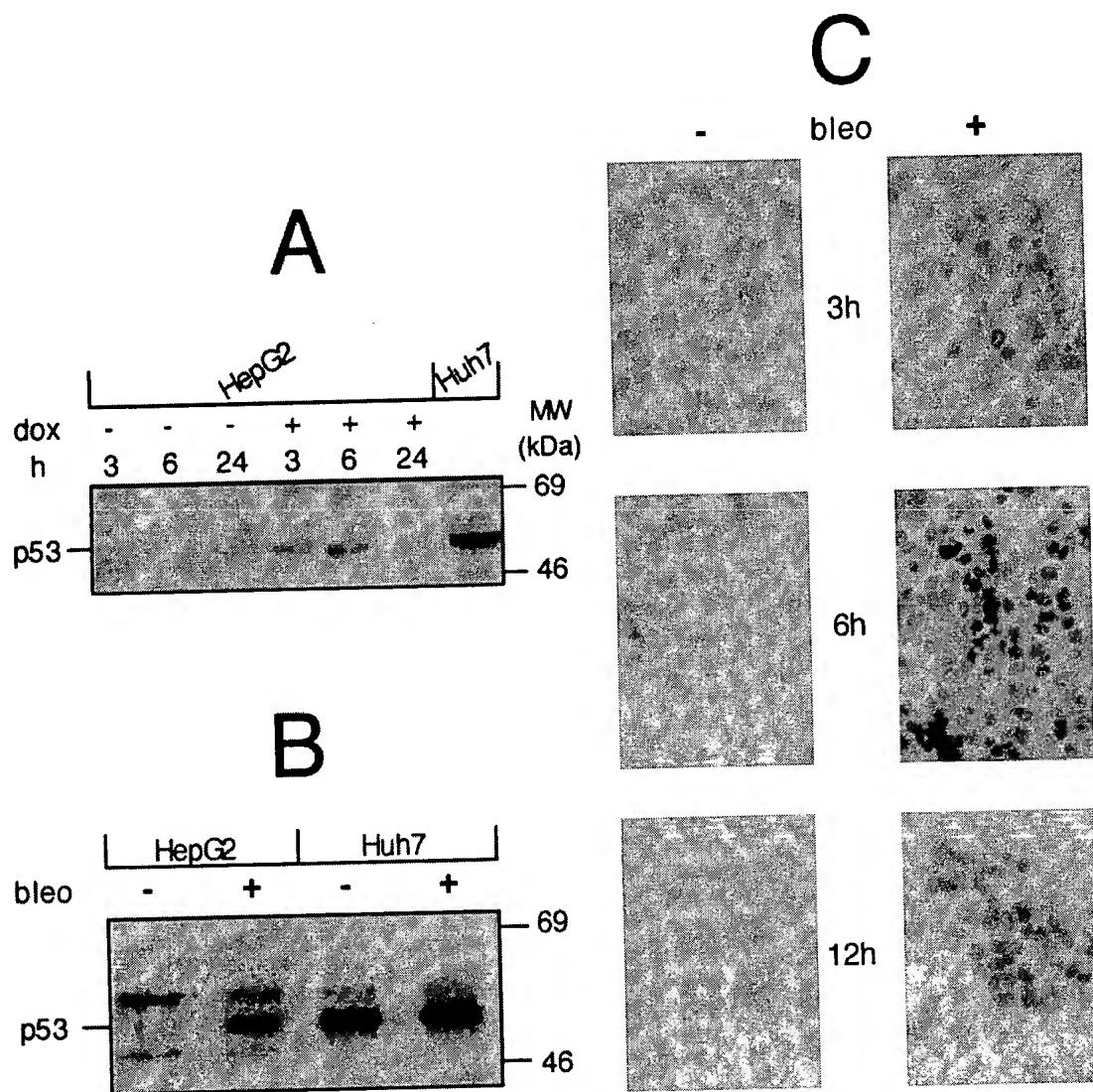


Figure 1. P53 induction in HepG2 cells after treatment with DNA-damaging agents. (A) Western blot of HepG2 cells treated with doxorubicin, 4 ng/ml (dox +), demonstrating a transient increase in p53 levels 3–6 h after the onset of treatment. Huh7 cells with mutant p53 served as positive control. (B) Autoradiography after endogenous labeling with [³⁵S]methionine and immunoprecipitation of p53 3–6 h after the onset of bleomycin treatment, 6 mg/ml (bleo +), demonstrates synthesis of p53. (C) Immunohistochemistry of HepG2 cells after treatment with bleomycin, 6 mg/ml (bleo +, right panel), reveals nuclear accumulation of p53. (Left panel) Untreated controls.

Digoxigenin-labeled RNA was detected immunologically by a polyclonal anti-digoxigenin sheep antibody conjugated with alkaline phosphatase (Boehringer), visualized using NBT and BCIP as substrate at room temperature overnight in the dark. Slides were mounted with Elvanol and viewed under a microscope (Zeiss, Axio-phot, Göttingen, Germany).

Detection of CD95 ligand mRNA expression by PCR

Total cellular RNA was prepared from 1×10^6 HepG2, Hep3B and Huh7 cells treated with 0.6 mg/ml and 6 mg/ml bleomycin for 24 h, using the Rneasy-Kit (Qiagen, Hilden, Germany), following the manufacturer's instructions. The efficiency of the RT and the amount of RNA used in the RT-PCR were verified by detection of the human β -actin mRNA with RT-PCR. 1 μ g of total cellular RNA was retrotranscribed after heat denaturation (3 min, 60°C) and annealing with oligo (dT) primers (16-mer, Perkin Elmer, Weiterstadt, Germany) in the presence of 75 U MnLV RT (Perkin Elmer), 67 μ M MgCl₂ and 63 μ M of each dNTP in 20 μ l for 45 min at 42°C. Reactions were stopped by heat inactivation for 5 min at 90°C. Aliquots of 10 μ l of the cDNA were then amplified in a DNA thermocycler (Stratagene, Heidelberg, Germany) with 2.5 U of Ampli Taq DNA-polymerase AS (Perkin Elmer), 10 pM of both upstream and downstream APO-1L Primers, and 2.5 μ M of both upstream and downstream human β -actin primers in a volume of 50 μ l. Each PCR cycle

consisted of a denaturation step (94°C, 1 min), an annealing step (56°C, 1 min), and an elongation step (72°C, 1 min). The primers used for amplification of the CD95L have been described recently (47). The PCR products were analyzed on a 1.4% TBE agarose gel.

Cytotoxicity assay

Crystal violet assay. A 100 μ l suspension of 7×10^3 cells was added to each well of 96-well plates 24 h before the assay. Various concentrations of anti-APO-1, bleomycin and F(ab)'₂-anti-APO-1 fragments were added to the target cells. After 3–72 h, cells were stained with 0.5% crystal violet in 20% methanol. Cell viability was determined by eluting the dye from the stained cells with 0.1 M sodium citrate/0.1 M citric acid and 50% ethanol, and absorbance was measured at 540 nm (48).

MTT assay. The MTT assay is a colorimetric assay based on the ability of the viable cells to reduce a soluble yellow tetrazolium salt (MTT) to blue formazan crystals. The cells were plated as described above for the crystal violet assay and appropriate concentrations of the different drugs were added. After 3–72 h 5 mg/ml MTT dye was added and the plates were incubated for 12 h. Optical density was determined by eluting the dye with isopropanol/formic acid, and absorbance was measured at 540 nm.

Crystal violet and MTT assay were performed in parallel. Results are shown as the mean of six independent wells \pm SD. To rule out that drug treated cells may simply arrest growth, making it difficult to de-

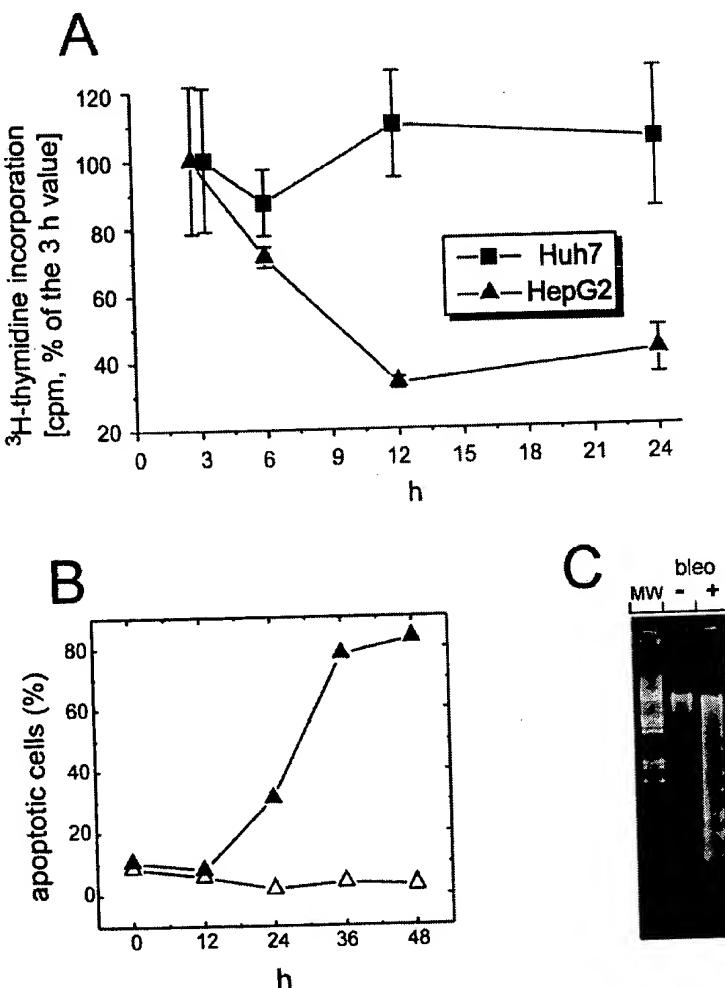


Figure 2. Induction of growth arrest and apoptosis in hepatoma cells by treatment with DNA damaging agents. (A) DNA-synthesis as measured by [³H]thymidine incorporation in HepG2 cells bearing wt p53 (\blacktriangle) and in Huh7 cells with mutant p53 (\blacksquare) treated with bleomycin, 6 mg/ml. Data are shown as the mean \pm SD of six independent culture plates and are expressed as percentage of untreated controls. (B) Kinetics of induction of apoptosis as detected by propidium iodide staining and FACS® analysis of HepG2 cells treated with bleomycin, 6 mg/ml (\blacktriangle). Untreated controls (\triangle). (C) Internucleosomal DNA fragmentation (DNA-“laddering”) after treatment of HepG2 cells with 6 mg/ml bleomycin (bleo +).

determine what percentage of the reduced crystal violet/MTT is due to apoptosis, all cytotoxicity assays were verified by FACS® analysis using the Nicoletti method (43) to assess specific apoptosis.

Results

Transient nuclear p53 accumulation after treatment of hepatoma cells with DNA damaging agents. Untreated HepG2 cells only showed basal p53 expression. Treatment of HepG2 cells with bleomycin or doxorubicin resulted in transient accumulation of p53 as detected by Western blotting (Fig. 1 A). An increased p53 signal was detectable 3 h after the onset of drug treatment. Maximum levels of about 5 to 10 times the control values were reached within 3 to 6 h after treatment. Elevated p53 levels returned to control values within 24 h after treatment. (Fig. 1 A). Likewise, 3 to 6 h after the onset of drug treatment of HepG2 cells a transient increase of [³⁵S]methionine-labeled p53 was observed (Fig. 1 B). In contrast to HepG2 cells, Huh7 cells expressing high constitutive levels of mutated p53 (p53-220 cys) (Fig. 1, A and B) did not show a significant increase in p53 expression after drug treatment (Fig. 1 B). Hep3B cells without endogenous p53 expression did not show a p53 signal by Western blotting before and after bleomycin treatment (data not shown). In HepG2 cells immunohistological analysis revealed increased p53 levels and showed a clear nuclear accumulation of the protein induced by bleomycin treatment (Fig. 1 C).

Induction of growth arrest and apoptosis by DNA damaging agents in human hepatoma cells. [³H]Thymidine incorporation in HepG2 cells after treatment with DNA damaging

agents showed a significant reduction in DNA synthesis. In contrast, Huh7 cells were remarkably insensitive to doxorubicin- and bleomycin treatment (Fig. 2 A).

We further investigated whether growth arrest after treatment by cytotoxic drugs was due to induction of apoptosis. Propidium iodide staining and FACS® analysis revealed induction of apoptosis in bleomycin-treated HepG2 cells (Fig. 2 B). Apoptosis was also shown by the typical "ladder" pattern of DNA fragmentation (Fig. 2 C).

Upregulation of the CD95 receptor in HepG2 cells and increased responsiveness towards induction of apoptosis by anti-APO-1 antibodies after bleomycin treatment. To study the involvement of the CD95 receptor/ligand system in drug-induced apoptosis we investigated CD95 mRNA and protein expression by semi-quantitative RT-PCR and FACS® analysis.

Bleomycin treatment resulted in enhanced transcriptional activation shown by augmentation of CD95 mRNA expression in HepG2 cells with wt p53. Huh7 cells (mut 220) and Hep3B cells (p53 *-/-*) did not upregulate CD95 mRNA upon bleomycin treatment (Fig. 3 A).

HepG2 cells showed a basal expression of the CD95 receptor in 10 to 15%, Huh7 cells in up to 3% and Hep3B cells in up to 3% of the cells. HepG2 cells revealed a marked upregulation of the CD95 receptor after drug treatment, up to 80% receptor expressing cells. This upregulation was only seen in wt p53 expressing HepG2 cells (Fig. 3 B). In contrast, neither Huh7 cells, bearing a point mutation in codon 220 of the p53 gene, nor Hep3B cells, lacking p53, responded to bleomycin treatment with upregulation of the CD95 receptor (Fig. 3 B).

Upregulation of CD95 receptor expression after drug treat-

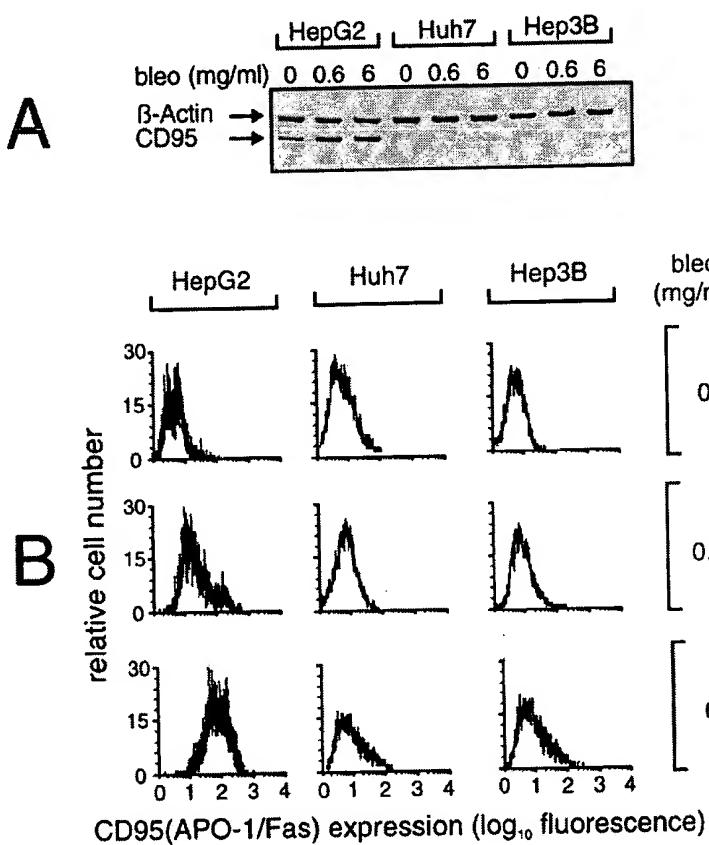
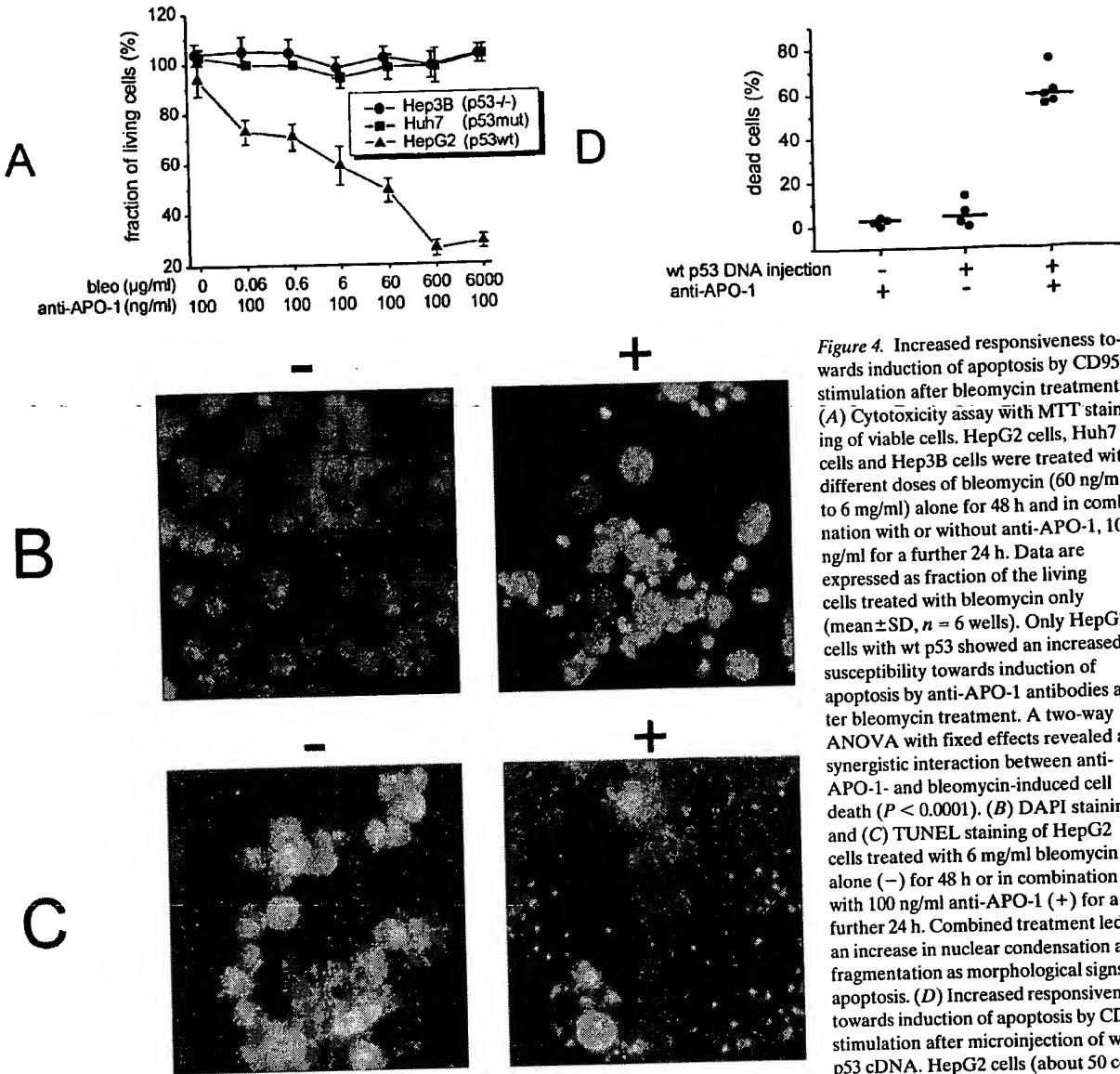


Figure 3. Upregulation of the CD95 receptor after bleomycin treatment. (A) Semi-quantitative RT-PCR analysis of CD95 receptor mRNA expression in HepG2, Hep3B and Huh7 cells after treatment with bleomycin 0.6 and 6 mg/ml. CD95 mRNA was induced in HepG2 cells (p53 wt) but not in Huh7 cells with mutant p53 and not in Hep3B cells without p53. (B) CD95 receptor expression by FACS® analysis in HepG2 with wt p53 (left column), Huh7 with mutant p53 (middle column) and Hep3B lacking p53 (right column). Cells were left untreated (upper panel) or received treatment with 0.6 mg/ml bleomycin (middle panel) or 6 mg/ml bleomycin (lower panel). Only HepG2 cells displayed enhanced CD95 receptor expression up to over 80% positive cells in response to bleomycin treatment in a dose dependent manner.



per experiment) were microinjected with wt p53 cDNA in 4 independent experiments. After 48 h 100 ng anti-APO-1 was added for 12 h and the surviving cells were counted under the microscope. Each dot represents the percentage of dead cells/microinjected cells from an independent well.

ment resulted in increased responsiveness towards CD95 stimulation. Treatment with anti-APO-1 alone led to specific cell death in up to 10% of the HepG2 cells, whereas it did not affect viability in Huh7 and Hep3B cells (Fig. 4A). In HepG2 cells treated with bleomycin for 48 h, CD95 receptor stimulation with anti-APO-1 resulted in induction of apoptosis in up to 75% of the cells, dependent on the concentration of bleomycin applied (Fig. 4, A-C). Stimulation of CD95 by anti-APO-1 and concurrent bleomycin treatment did not only show an additive effect but synergized in the induction of apoptosis in p53 wt hepatoma cells. To test for synergy a two-way ANOVA model with fixed effects was performed ($P < 0.0001$).

As a control, the p53 mutant Huh7 cells and the p53 null Hep3B cells were tested. Anti-APO-1 did not produce addi-

tional toxicity in comparison with bleomycin treatment alone in Huh7 and in Hep3B cells (Fig. 4A). Similar as bleomycin treatment microinjection of wt p53 cDNA into HepG2 cells resulted in nuclear accumulation of p53 (not shown) and increased sensitivity towards CD95 mediated apoptosis (Fig. 4D).

Induction of apoptosis by bleomycin is mediated by increased expression of CD95L. To obtain further insight into the mechanism of bleomycin-induced apoptosis, we also analyzed CD95L expression upon drug treatment. Fig. 5A shows by *in situ*-hybridization that CD95L mRNA is highly elevated upon treatment of HepG2 cells with bleomycin. This was confirmed by PCR analysis. Bleomycin, at concentrations measured in patients' sera during therapy, stimulated CD95L mRNA expression *in vitro* in HepG2 cells. Hep3B cells (p53^{-/-}) did

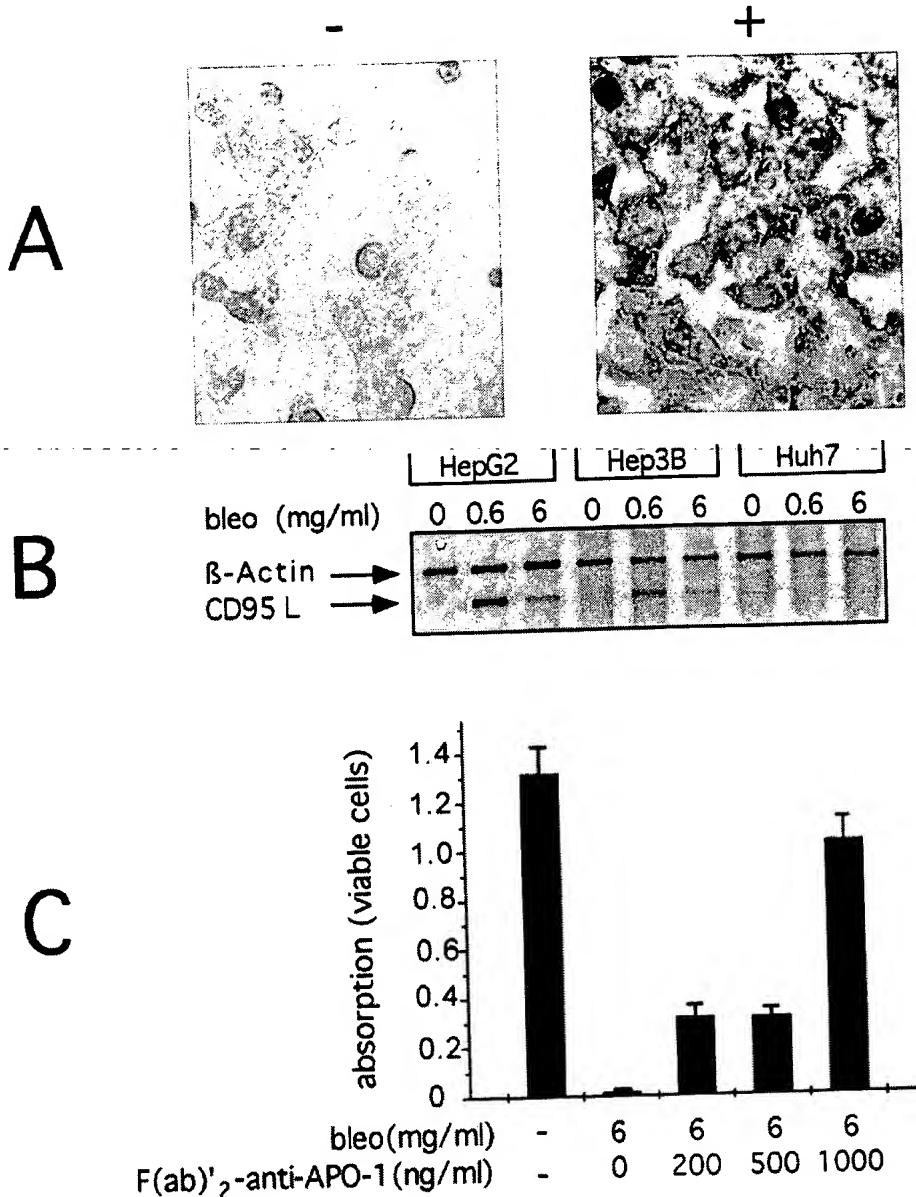


Figure 5. Induction of the CD95L by bleomycin. (A) CD95L mRNA in-situ hybridization of untreated HepG2 cells (-) and after treatment with 6 mg/ml bleomycin for 24 h (+). (B) Semi-quantitative PCR analysis of CD95L mRNA expression in HepG2, Hep3B and Huh7 cells upon treatment with bleomycin 0.6 mg/ml and 6 mg/ml. CD95L mRNA expression was induced in HepG2 cells (p53wt) and in Hep3B cells (p53-/-) but not in Huh7 cells (p53mut). (C) Inhibition of bleomycin-induced apoptosis by F(ab)'2-anti-APO-1. Cytotoxicity assay with crystal violet staining of viable cells. HepG2 cells were treated with 6 mg/ml bleomycin for 72 h alone or in combination with CD95 receptor-antagonizing F(ab)'2-anti-APO-1 antibody fragments. F(ab)'2-anti-APO-1 antibody fragments were able to preserve up to 80% of the cells from death.

also respond by CD95L upregulation towards bleomycin treatment. In contrast, bleomycin treatment of Huh7 with mutant p53 did not result in upregulation of the CD95L (Fig. 5 B). These findings suggested that bleomycin-induced apoptosis in HepG2 cells with wt p53 might be mediated, at least in part, by the activity of the CD95L. The assumption that apoptosis induced by bleomycin in HepG2 cells is mediated via the CD95/CD95L system was strongly supported by CD95 blocking experiments. Fig. 5 C shows that bleomycin-induced apoptosis after 72 h of culture can almost completely be inhibited by F(ab)'2-anti-APO-1 antibody fragments, known to interfere with CD95 receptor/ligand interaction (37). Thus, bleomycin-induced apoptosis is mediated, at least in part, through a death signal by increased expression of CD95L binding to the CD95 receptor. Furthermore, in long-term [³H]thymidine-labeled cultures, we could show that treatment with F(ab)'2-anti-APO-1

antibody fragments affects true survival of the cells. Thus, the proliferation of treated HepG2 cells was double to triple that of HepG2 cells treated with bleomycin only. These data show that F(ab)'2-anti-APO-1 antibody treatment affects true survival in HepG2 cells treated with cytotoxic drugs and does not simply delay the kinetics of death in the cells.

Anticancer drugs with different mechanisms of action induce apoptosis via the CD95 system. Cisplatin and methotrexate have been shown to trigger nuclear accumulation of the tumor suppressor protein p53 (49). Cisplatin and methotrexate treatment lead to a marked increase in CD95 mRNA and protein expression in wild type p53 expressing HepG2 cells. Upon cisplatin treatment, CD95 expression by FACS® analysis was enhanced up to 50% positive cells. Methotrexate treatment resulted in CD95 receptor upregulation up to 70% positive cells. As upon bleomycin treatment neither Huh7 cells (mut 220)

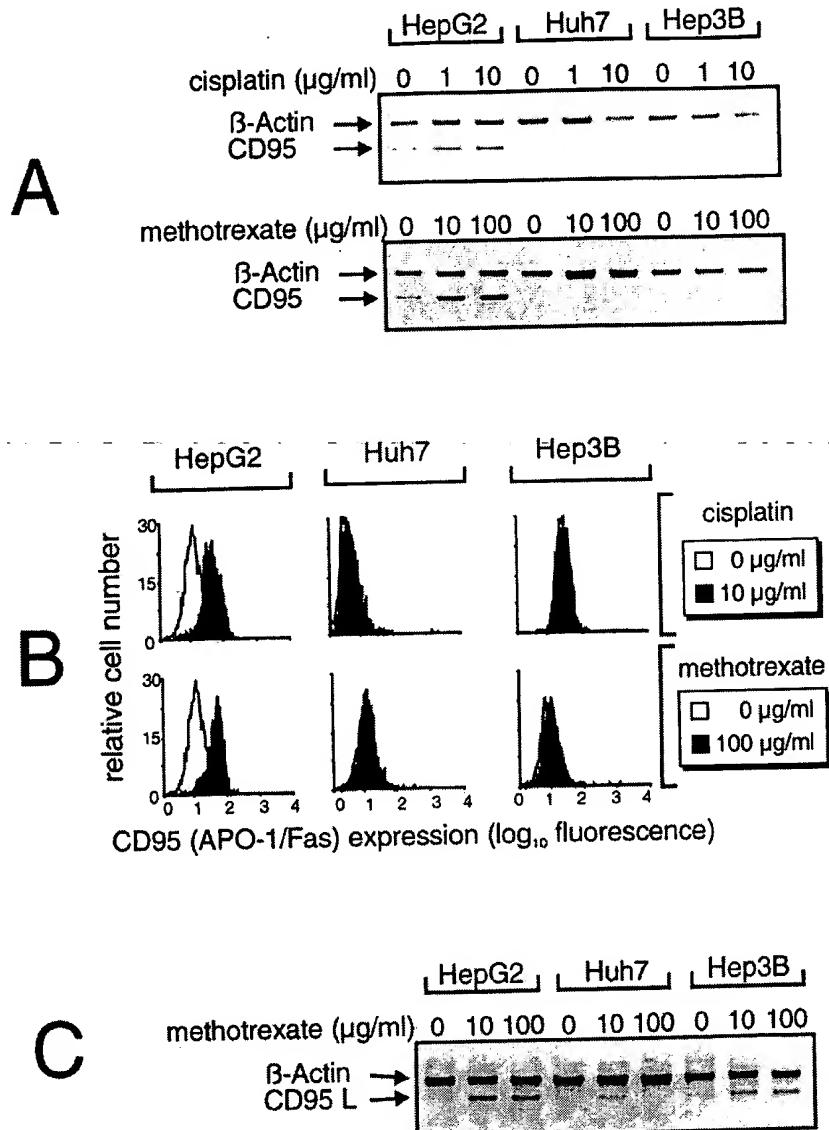


Figure 6. Induction of the CD95 system by anticancer drugs with different mechanisms of action. (A) Semi-quantitative PCR analysis of CD95 receptor mRNA expression in HepG2, Huh7 and Hep3B cells. CD95 mRNA was induced in HepG2 cells (p53 wt) by cisplatin at 1 µg/ml and 10 µg/ml and by methotrexate at 10 µg/ml and 100 µg/ml. Huh7 cells with mutant p53 and p53 deficient Hep3B cells did not respond to anticancer therapy with upregulation of CD95 receptor mRNA. (B) CD95 receptor expression by FACS® analysis in HepG2 with wt p53 (left column), Huh7 with mutant p53 (middle column) and Hep3B lacking p53 (right column). Only HepG2 cells displayed enhanced CD95 receptor expression up to 50% positive cells in response to cisplatin and up to 70% positive cells in response to methotrexate. (C) CD95 ligand expression after treatment with methotrexate. Semi-quantitative PCR analysis of CD95 ligand mRNA expression in HepG2, Hep3B, and Huh7 cells after treatment with methotrexate at 10 µg/ml and 100 µg/ml. Methotrexate induced CD95 ligand mRNA expression in all three cell lines with the exception of the high dose methotrexate in Huh7. The CD95 mRNA expression was weaker in Huh7 cells with mutated p53 than in p53 wt and p53 deficient cells.

nor Hep3B cells ($p53^{-/-}$) upregulated CD95 mRNA and protein in response to anticancer therapy (Fig. 6, A and B). This was confirmed by stimulation of CD95 by anti-APO-1 antibodies and concurrent cisplatin or methotrexate treatment. In HepG2 cells treated with cisplatin or methotrexate for 48 h, CD95 receptor stimulation with anti-APO-1 antibodies resulted in further induction of apoptosis in 50% of the cells compared to cisplatin and methotrexate treatment alone. As in combined bleomycin and anti-APO-1-treatment, anti-APO-1 antibodies did not further enhance apoptosis in comparison with cisplatin and methotrexate alone in Huh7 and in Hep3B cells (data not shown).

Methotrexate treatment induced CD95L mRNA expression at 10 µg/ml and 100 µg/ml in all three cell lines, with the exception of the high dose methotrexate in Huh7 cells. CD95L mRNA expression was weaker in Huh7 cells with mutant p53 than in HepG2 cells (wt p53) and in Hep3B cells ($p53^{-/-}$) (Fig. 6 C). Cisplatin treatment was not found to initiate CD95L

mRNA expression at 12 h, 24 h and 48 h after the onset of treatment (data not shown).

Thus, methotrexate- and cisplatin-induced apoptosis involves similar mechanisms as shown for bleomycin treatment.

Discussion

The data in this paper show that cytotoxicity in hepatoma cells by the anti-cancer drugs bleomycin, doxorubicin, cisplatin, and methotrexate is mediated by induction of apoptosis. In addition, bleomycin treatment resulted in accumulation of nuclear p53 and stimulated increased expression of cell surface CD95 in the drug-sensitive wt p53 hepatocellular carcinoma cell line HepG2. Bleomycin did not increase CD95 in two hepatoma lines with dysfunctional p53. Bleomycin enhanced CD95-induced apoptosis in HepG2 cells. In contrast, in p53 mutant Huh7 cells and in p53 negative Hep3B cells, stimulation of

CD95 does not increase death after bleomycin treatment. Furthermore, bleomycin treatment increased expression of CD95L mRNA. Finally, bleomycin-induced apoptosis in HepG2 cells could almost completely be inhibited by antibodies which block binding of CD95L to the CD95 receptor. Thus, bleomycin-induced apoptosis was shown to be mediated by stimulation of the CD95 receptor/ligand system.

Upregulation of the CD95 system might therefore be a major cause of apoptosis in anticancer therapy. We could show that anticancer drugs with different mechanisms of action, like cisplatin and methotrexate, do also exert their effects via the CD95 (APO-1/Fas) receptor/ligand system. Subsequent to the investigation of CD95 receptor and ligand expression in hepatoma cells, we are currently investigating the involvement of the CD95 receptor/ligand system in the chemotherapeutic treatment of different solid cancers, including gastrointestinal and breast cancer cell lines. Preliminary data indicate that activation of the CD95 system is involved in induction of apoptosis in response to anticancer therapy in different solid human tumor cell lines. Furthermore, data by Friesen et al. (50) show CD95L upregulation in drug induced apoptosis in leukemia cell lines. Apoptosis by the CD95 system might therefore play a general role for cytotoxicity by anti-cancer drugs in a variety of cells of different histotype.

In addition to the data of Friesen et al. (50), which show upregulation of the CD95L in leukemia cell lines, constitutively expressing CD95, we could demonstrate that anticancer agents at concentrations relevant for therapy of patients lead to an upregulation of both, the CD95 receptor and the CD95 ligand in cell lines of a solid human tumor. Thus, upregulation of the CD95 receptor and/or the CD95 ligand might render a tumor cell sensitive towards chemotherapy. On the contrary, drug resistance could result from downregulation of the CD95 receptor. Thus, anticancer agents, stimulating CD95L expression cannot induce apoptosis in tumor cells that have downregulated CD95. This is supported by our findings in Hep3B cells which show stable CD95 receptor expression of only 3%. In these cells CD95 ligand induction by cytotoxic agents does not induce further toxicity. Indeed, in studying the CD95 system in patients with HCC, we could demonstrate loss of CD95 receptor and gain of CD95L expression compared to normal liver tissue (Strand, S., P.H. Krammer and P.R. Galle, manuscript submitted for publication). Alterations of the CD95 receptor/ligand system might constitute new mechanisms of drug resistance. These mechanisms might exist apart from the known mechanisms of amplification of mdrl, overexpression of bcl-2, BCR-ABL, or mdm2 proteins, or underexpression of DNA topoisomerase II.

Chemoresistance has also been linked to mutations in p53. This is supported by our finding of a reduced responsiveness of Huh7 cells bearing mutant p53 towards bleomycin treatment as compared to the responsiveness of HepG2 cells with wt p53. This demonstrates the requirement of wt p53 for chemotherapy-induced apoptosis in these cells. This requirement has also been shown for a variety of other non-hepatoma cell lines and in animal experiments (3, 51). However, there are also conflicting data suggesting that the relation between p53 status and chemoresistance may be tissue-specific (52-54). In fact, mutations of p53 in many human malignancies have been linked to poorer prognosis (55, 56).

The action of p53 and its effect on the CD95 receptor/ligand system is unclear. Our data show that stimulation of the

CD95 receptor did only occur in HepG2 cells with intact p53. Hepatoma cells with mutant p53 and p53 null cell lines did not upregulate the CD95 receptor after cytotoxic therapy. This is supported by our data from the microinjection experiments. Direct injection of wt p53 cDNA into the nucleus of HepG2 cells increased susceptibility towards CD95 mediated apoptosis. These data are in accordance with data by Owen-Schaub et al. (16) demonstrating, that only the wt form of a temperature sensitive mutant of p53 could induce CD95 expression. Stimulation of CD95 expression by p53 was shown to be dependent on transcriptional activity (16). It is possible therefore, that the promoter of the CD95 receptor gene is stimulated by p53 dependent transcriptional activity directly. Alternatively, p53 might activate transcription factors which regulate the CD95 receptor gene. Intact wt p53 is required to mediate such activity. Thus, wt p53 might sense DNA damage in cells treated with anti-cancer drugs such as bleomycin and stimulate expression of the CD95 receptor gene. Our laboratory is currently undertaking experiments to further investigate a possible p53 effect on the CD95 receptor promoter.

Upregulation of the CD95 ligand after bleomycin treatment was demonstrated in p53 wt and p53 deficient hepatoma cell lines, but not in cell lines with mutant p53. This would argue for a gain of function of mutant p53 in the sense of a possible inhibition of CD95L expression. However, methotrexate treatment resulted in a slight upregulation of the CD95L in Huh7 cells. Thus, the regulation of the CD95L does not solely depend on p53, and p53 independent mechanisms must be involved in the regulation of CD95L.

The function of the CD95 system is complex and involves expression of membrane-bound or soluble CD95 receptor or ligand, a sensitive CD95 receptor signaling pathway and a balance of pro-apoptotic and anti-apoptotic programs (57). Recently, part of the CD95 signaling cascade has been elucidated (reviewed in references 58 and 59). Oligomerization of the CD95 receptor by agonistic anti-CD95 antibodies or by CD95L is essential for the CD95-mediated apoptotic signal. Oligomerization of CD95 causes binding of FADD and the effector protease FLICE, a novel member of the ICE/CED-3 family to the intracellular CD95 associated death-inducing signaling complex (DISC) (60). The assembly of this signaling complex occurs in a hierarchical manner upon receptor activation. The death domain of the receptor binds to the death domain of the adapter molecule FADD, which binds and activates FLICE to generate the most apical enzymatic component of a death cascade composed of other ICE/CED-3 family members (61). Further downstream, after DISC formation, ICE-like proteases (62, 63) propagate the apoptotic signal and catalyze a variety of substrates. The CD95 signaling cascade is still incomplete and might be positively or negatively modulated by members of the bcl-2 family (64, 65) or the activity of kinases or phosphatases such as RIP, HLP, or FAP-1 (66). Moreover, members of the bcl-2 family have recently been shown to interfere with both the p53 and the CD95 system in some cells. It has been shown that bcl-2 overexpressing mice do not respond to CD95 stimulation with liver failure, in contrast to wt-controls (67). On the other hand, bcl-2 and its antagonist bax have been shown to be regulated by p53 (68, 69).

Drug sensitivity or resistance will be affected by alterations of any components of this pathway. Insight into the interactions of the components of the CD95 signaling pathway will therefore provide the basis for modulation of CD95 sensitivity

and for the development of drugs which either specifically enhance or reduce apoptosis induced by anti-cancer drugs.

It is likely that these data have profound implications for the future management of drug treatment in patients with malignant diseases, particularly for the understanding of drug sensitivity and resistance. Sensitivity to anticancer therapy will therefore depend, at least in part, on the integrity of the p53 and the CD95 receptor/ligand system. In contrast, malfunction of these systems might result in development of drug resistance. This might be useful in a clinical setting to increase cytostatic anti-tumor action and to reduce systemic side effects.

These data may also have implications for p53 gene therapy. In the first phase I clinical trial of p53 gene therapy in lung cancer, Roth and colleagues (70) demonstrated that introduction of wild type p53 into tumor cells resulted in induction of apoptosis of the tumor cells and in control of tumor growth in patients with relapsed lung cancer (whose tumors all carried a p53 mutation). Our data suggest, that upregulation of the CD95 system may account for the apoptosis inducing effect of wild type p53. Thus, wt p53 gene transfer and chemotherapy might cooperate to optimize future cancer therapy.

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